

CHANNEL BLOCKING COMPOUNDS

TECHNICAL FIELD

The present invention relates to ion channel blocking compounds in particular, compositions containing alkaloid compounds termed lolitrems for use as ion
5 channel antagonists and to methods and uses of these compositions. More specifically, the lolitrem compounds are derived from the species *Neotyphodium* (formally *Acremonium*) *lolii* and are used as potassium channel antagonists.

BACKGROUND ART

10 Ion channels are defined as transmembrane pores that present a central aqueous pore that can be opened by conformational change to allow ions to cross a lipid bilayer down their electrochemical gradients. Some degree of ion specificity is usually observed and typically a million ions per second flow through the channel. Channels may open spontaneously, like the potassium leak channel, or they may
15 be voltage-gated, like the voltage-gated sodium channel or ligand-gated, like the acetylcholine receptor.

Ion channels generally are the subject of much research to understand the roles they have in normal physiological systems and in disease states.

Potassium ion channels are selective for potassium ions. There are diverse types
20 of potassium ion channels with different functions, for example: voltage-gated potassium channels, delayed rectifier channels, M channels, A channels, inward rectifier channels, and calcium-activated potassium channels.

Large or high conductance calcium-activated potassium channels are also termed as BK channels, K_{Ca} , maxi-K. *Slowpoke* (*Slo*) is the name of the gene that encodes

the pore-forming α subunit of the channel, e.g. *hSlo* – human gene, *mSlo* – mouse, *dSlo* drosophila – fruit fly. Accessory β subunits (β_1 - β_4) associate with the α subunit to generate BK channel diversity. BK channels are gated by Ca^{++} and membrane potential with a unit conductance of 100 to 300 picoSiemens (pS)

- 5 Calcium-activated potassium channels also include intermediate conductance (IK) and small conductance (SK) channels. IK potassium channels are more sensitive to Ca^{++} than BK channels and are gated only by internal Ca^{++} ions, having a unit conductance of 25 to 100 pS. SK channels are also highly sensitive to Ca^{++} and have minimal voltage sensitivity, and a unit conductance of 2 to 25 pS.
- 10 For the purposes of this specification, the term 'BK channel' or 'potassium channel' or similar variations will be referred to although, this should not be seen as limiting.

BK channels are expressed in many tissues, including muscle and brain and regulate important physiological functions (for review see Gribkoff et al., 2001a). They have a role in regulation of blood pressure and are implicated in hypertension

15 (Brenner et al., 2000a, Amberg and Santana, 2003, Amberg et al., 2003). They are activated in response to depolarising voltages and to increased intracellular calcium. Their activation results in efflux of potassium ions causing hyperpolarisation which dampens cellular excitability. BK channels are expressed in most tissues and control a large variety of physiological processes including

20 smooth muscle tone, neurosecretion and hearing.

In blood vessels, BK channels oppose vasoconstriction, allowing vasorelaxation and thereby regulate arterial tone (i.e. blood pressure) (Nelson et al., 1995).

In the brain, they modulate action potential waveform, repetitive firing and neurotransmitter release (Shao et al., 1999, Golding et al., 1999, Hu et al., 2001).

25 They are also expressed in the cochlea of the ear where they have a specialised role in frequency tuning of hair cells, acting in concert with other ion channels

(Gribkoff, et al 2001a; Orio et al 2002).

BK channels are also expressed in other tissues where their role is not known, e.g. ovary, testis and kidney (Brenner et al., 2000b).

Compounds that block (inhibit) a biologic activity or process in this transfer of ions
5 across a cell membrane are called 'blockers'. They may also be termed
'antagonist compounds' as the compounds reduce or prevent ion transfer. For the
purposes of this specification the term 'antagonist' will be used. This should not,
however be interpreted as limiting.

The function of BK channels is modulated by a variety of compounds (for review
10 see Kaczorowski and Garcia, 1999, Kaczorowski et al., 1996).

Known marketed drugs that block potassium channels include Glyburide™,
Glipizide™ and Tolbutamide™. Other naturally occurring toxins that are known to
block potassium channels include Apamin, Iberiotoxin, Charybdotoxin, Noxiustoxin
and Kaliotoxin. US 5,541,208 describes uses of these blockers and the use of
15 paxilline, a further blocking compound, and is incorporated herein by reference.

Lolitrems compounds belong to the broader group of alkaloid compounds
incorporating indole diterpenes.

Assay techniques for identifying lolitrems compounds are known, for example see
NZ 236879.

20 Lolitrems compounds are present in perennial ryegrass (*Lolium perenne*) infected
with the endophytic fungus *Neotyphodium* (formally *Acremonium*) *lolii* (Lane et al.,
2000). Lolitrems compounds have been extracted from endophyte-infected
ryegrass seed (Gallagher et al., 1981, Miles et al., 1994, Munday-Finch et al.,
1995, Munday-Finch et al., 1996, Munday-Finch et al., 1997, Munday-Finch et al.,
25 1998).

Endophytes are symbiotic fungi and are prevalent in at least New Zealand pastures. The fungal metabolites from these endophytes are thought to serve as chemical defence systems for the fungi that produce them. They may also be of use in protecting the food source from consumption by other organisms (US 4,973,601).

The lolitrems are neurotoxic indole-diterpenes and are the principal causative agents of ryegrass staggers. This is a condition in which animals grazing on endophyte infected ryegrass-dominant pastures develop ataxia, tremors, and hypersensitivity to external stimuli. The lolitrem neurotoxin (staggers) reaction is long acting but is however completely reversible (Smith et al 1997, McLeay et al 1999). The time course of tremors induced by lolitrem B is dramatically different from that of other indole diterpenes, for example paxilline and analogues. When injected into mice, paxilline analogues induce tremors of rapid onset and short duration while tremors induced by lolitrem derivatives take hours to reach maximum intensity and last for days (Munday-Finch, 1997). Tremors induced by lolitrem are also longer in duration than those induced by the indole diterpene, aflatrem (Gallagher and Hawkes, 1996).

Whilst at least some lolitrems and other indole diterpenes, for example paxilline, are known to cause tremorgenicity (tremorgenic mycotoxins) there is no proven link between tremorgenicity and BK channel blocking.

Some linkage is inferred between tremorgenicity and neurotransmitter release (Mantle 1983, Gallagher et al 1986, Smith et al 1997, McLeay et al 1999, Wang et al 2003).

Some alkaloid compounds and more specifically indole diterpenes, block BK channels (e.g. paxilline, US 5,541,208) and some do not (McMillan et al., 2003). The alkaloids that inhibit BK channels include both tremorgens and non-

tremorgens. Structural moieties that are important for BK channel antagonism have been determined for some paxilline derivatives (Knaus et al 1994). However, for other types of indole diterpenes (e.g. lolitrems) whether a given compound will inhibit the BK channel cannot yet be predicted from structure alone but must be
5 determined empirically.

Within a given structural class of indole diterpene, tremorgenicity cannot be predicted by structure. For example, while paxilline and lolitrem B are tremorgenic, lolilline, which is intermediate in structure between the two, is non-tremorgenic.

The structural features required for tremorgenicity are also different for each group
10 of structurally related indole diterpenoid compounds. An acetal-linked isoprene unit, the presence of A/B rings and the stereochemistry at the A/B ring junction have been identified as important structural features for the tremorgenicity of lolitrem derivatives. Different structural features are required for tremorgenicity for other indole diterpene compounds.

15 As channel blockers have a variety of pharmaceutical uses and have been found to be beneficial for treatment of some diseases (for example Parkinson's disease, US 5,541,208), such blocking compounds are of interest, particularly in the development of new therapies. A BK channel modulator that opens channels has been investigated as a neuroprotective drug in treating ischemic stroke (Gribkoff et
20 al., 2001b).

It is an object of the present invention to provide an alternative ion channel blocking compound or at least to provide the public with a useful choice.

All references, including any patents or patent applications cited in this specification are hereby incorporated by reference. No admission is made that
25 any reference constitutes prior art. The discussion of the references states what their authors assert, and the applicant reserves the right to challenge the accuracy

and pertinency of the cited documents. It will be clearly understood that, although a number of prior art publications are referred to herein, this reference does not constitute an admission that any of these documents form part of the common general knowledge in the art, in New Zealand or in any other country.

- 5 It is acknowledged that the term 'comprise' may, under varying jurisdictions, be attributed with either an exclusive or an inclusive meaning. For the purpose of this specification, and unless otherwise noted, the term 'comprise' shall have an inclusive meaning - i.e. that it will be taken to mean an inclusion of not only the listed components it directly references, but also other non-specified components
- 10 or elements. This rationale will also be used when the term 'comprised' or 'comprising' is used in relation to one or more steps in a method or process.

Further aspects and advantages of the present invention will become apparent from the ensuing description which is given by way of example only.

15 **DISCLOSURE OF THE INVENTION**

It has been found by the inventors that lolitrem compounds are antagonists of potassium (BK) ion channels.

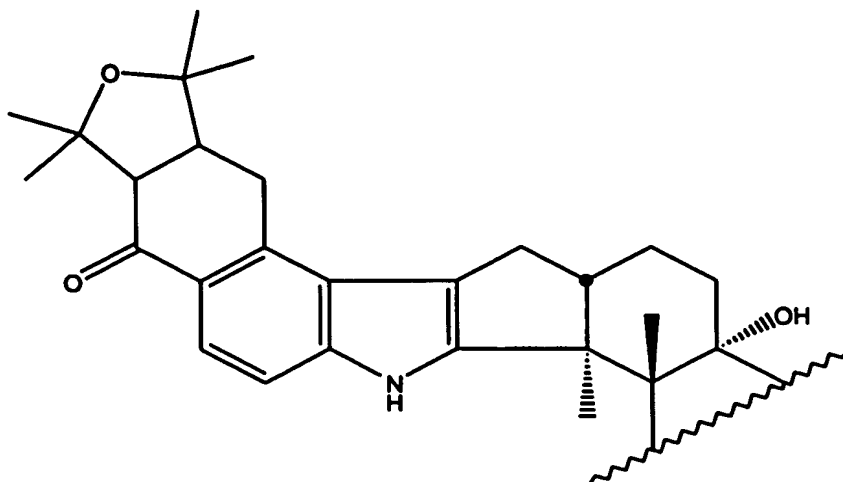
- For the purposes of this specification the term 'ion channel' refers to transmembrane pores that present a central aqueous pore that can be opened by
- 20 conformational change to allow ions to cross a lipid bilayer down their electrochemical gradients.

For the purposes of this specification the term 'antagonist' refers to compounds that reduce or prevent ion transfer across a cell membrane.

According to one aspect of the present invention there is provided a composition

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that contains a pharmacologically effective amount of at least one BK channel antagonist compound containing the moiety shown in structure (I):



STRUCTURE (I)

5 or derivatives thereof.

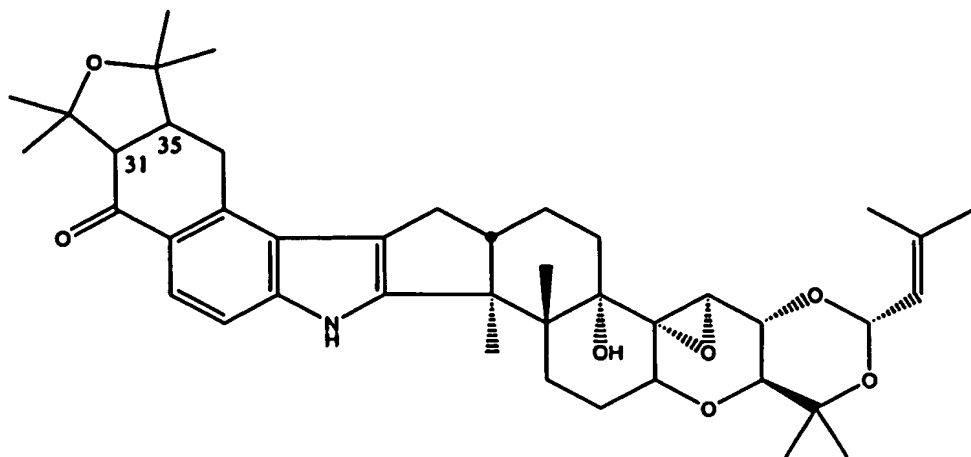
Preferably, derivatives of structure (I) are selected from the group consisting of: salts, analogues, isomers, and combinations thereof.

Preferably, the antagonist compound is selected from the group consisting of:

10 lolitrem B, lolitrem A, lolitrem F, 31-*epi*lolitrem F, 31-*epi*lolitrem B, lolitrem E, lolitrem E acetate, lolitrem L, lolitrem G, lolitrem C, lolitrem M, lolitriol, lolitriol acetate, lolitrem N, lolitrem J, lolitrem H, lolitrem K, lolicine A and B, 30-desoxy lolitrem B-30 α -ol, 30-desoxy-31-*epi*lolitrem B-30 α -ol, 30-desoxylolitrem B-30-ene lolilline and combinations thereof.

Preferably, the antagonist compound is structure (II):

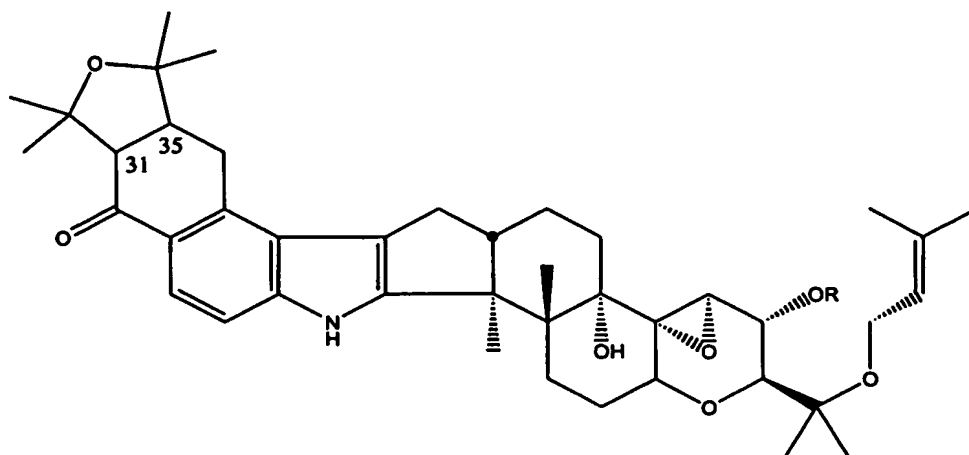
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STRUCTURE (II)

which includes compounds selected from the group consisting of: lolitrem B = 31 α ,
 35 β stereochemistry; 31-*epi*lolitrem B = 31 β , 35 β stereochemistry; lolitrem F =
 5 31 α , 35 α ; 31-*epi*lolitrem F = 31 β , 35 α .

Preferably, the antagonist compound is structure (III):

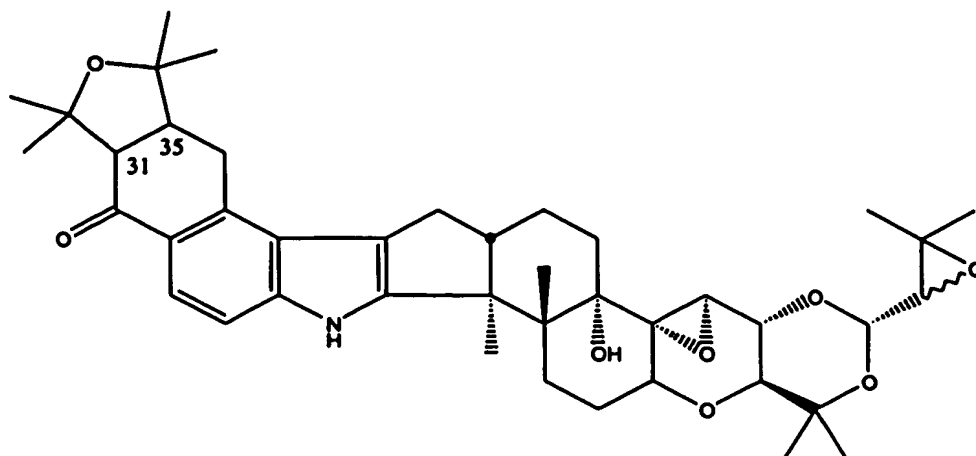


STRUCTURE (III)

which includes compounds selected from the group consisting of: lolitrem E = 31 α ,
 10 35 β stereochemistry where R = H or acetate; lolitrem L = 31 α , 35 α
 stereochemistry where R = H or acetate.

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Preferably, the antagonist compound is structure (IV):

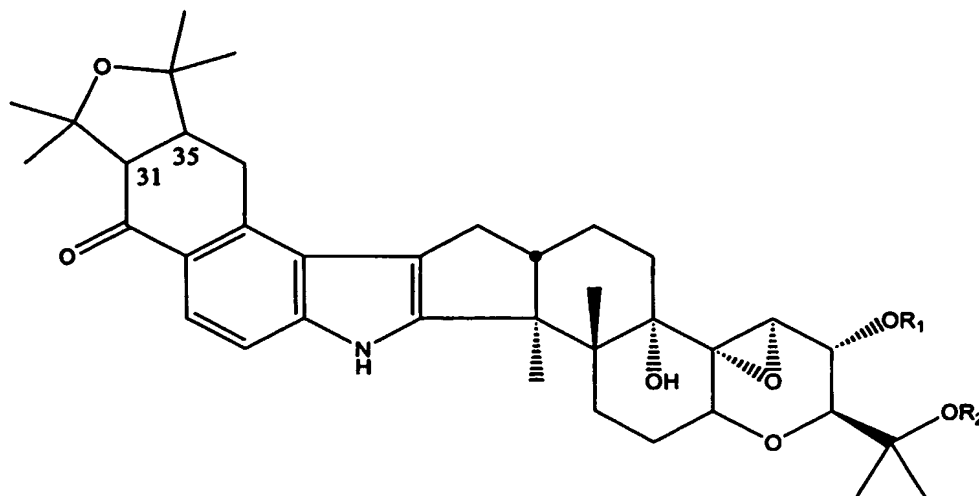


STRUCTURE (IV)

which includes compounds selected from the group consisting of: lolitrem A = 31 α ,

5 35 β stereochemistry; lolitrem G = 31 α , 35 α stereochemistry.

Preferably, the antagonist compound is structure (V):



STRUCTURE (V)

which includes compounds selected from the group consisting of: lolitriol; = 31 α ,

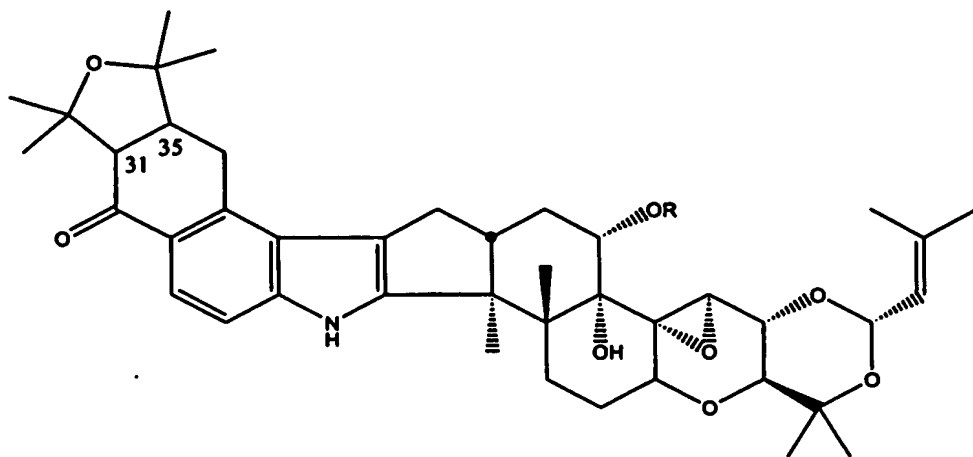
10 35 β stereochemistry where R₁ = H or acetate and R₂ = H; lolitrem N = 31 α , 35 α

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stereochemistry where $R_1 = \text{H}$ or acetate and $R_2 = \text{H}$; Lolitrem J = 31α , 35β

stereochemistry where $R_1 = \text{H}$ or acetate and $R_2 = \text{acetate}$.

Preferably, the antagonist compound is structure (VI):

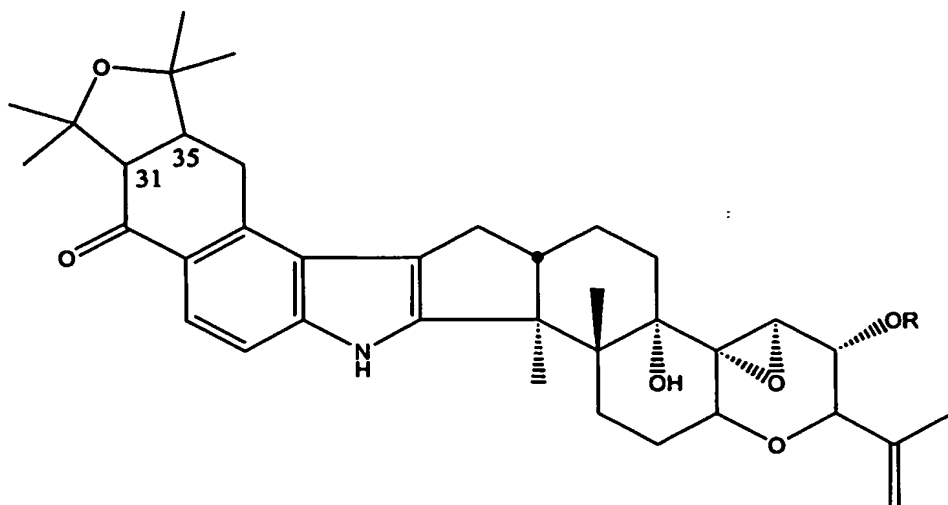


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STRUCTURE (VI)

which includes lolitrem H = 31α , 35β stereochemistry where $R = \text{H}$ or acetate.

Preferably, the antagonist compound is structure (VII):

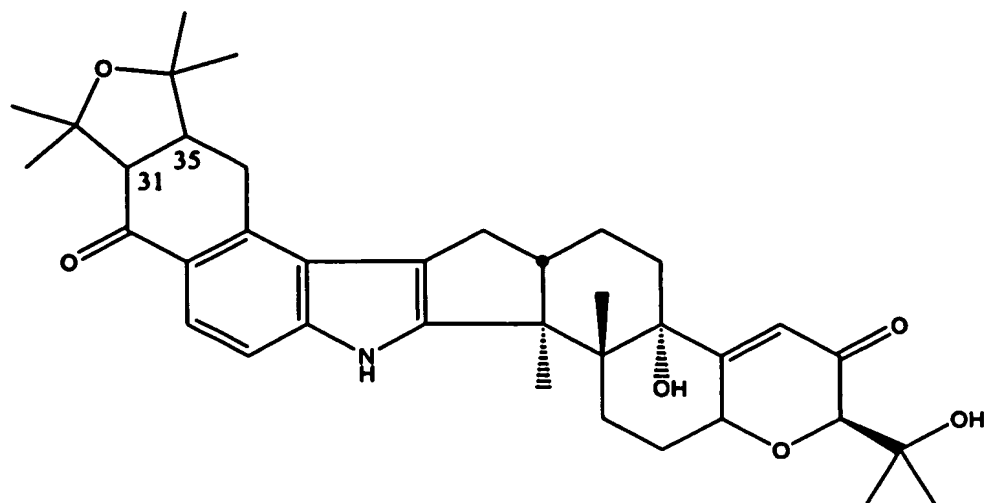


STRUCTURE (VII)

10 which includes lolitrem K = 31α , 35β stereochemistry, where $R = \text{H}$ or acetate.

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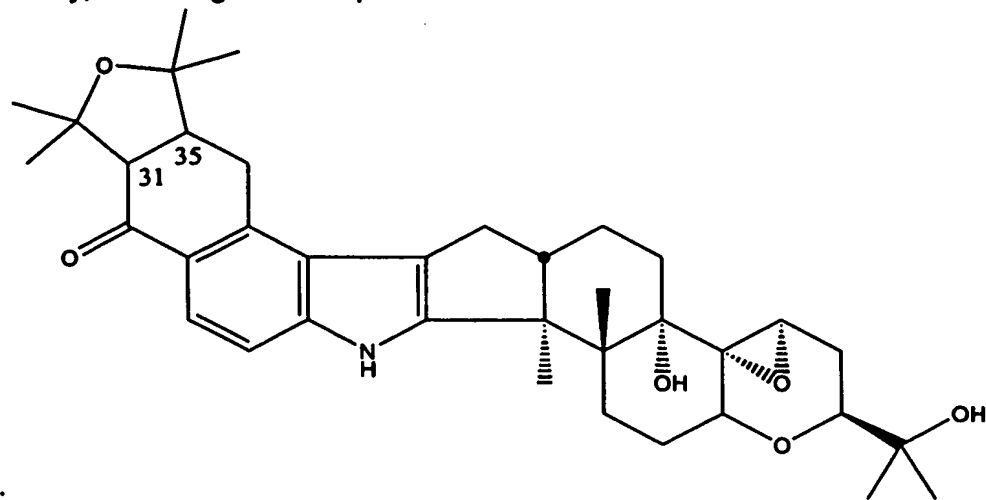
Preferably, the antagonist compound is structure (VIII):



STRUCTURE (VIII)

which includes lolilline = 31α , 35β stereochemistry.

5 Preferably, the antagonist compound is structure



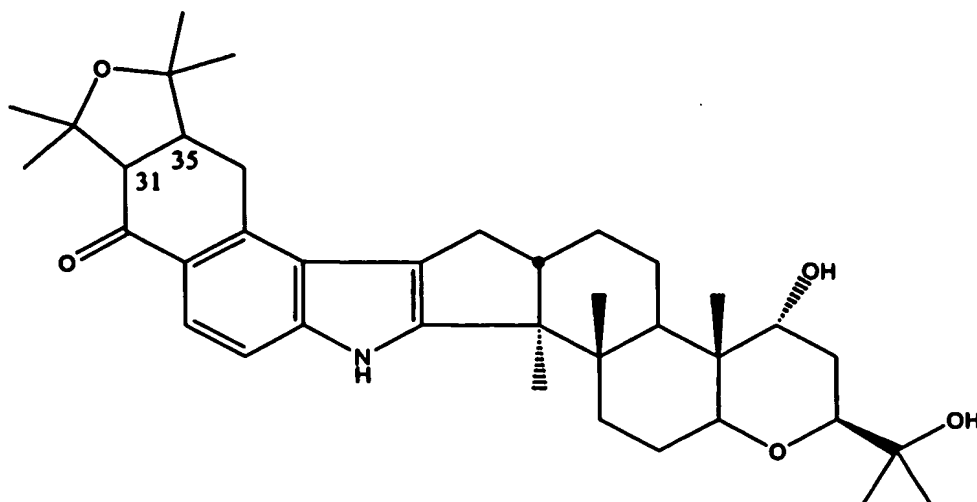
(IX):

STRUCTURE (IX)

which includes lolitrem M = 31α , 35β stereochemistry.

Preferably, the antagonist compound is structure (X):

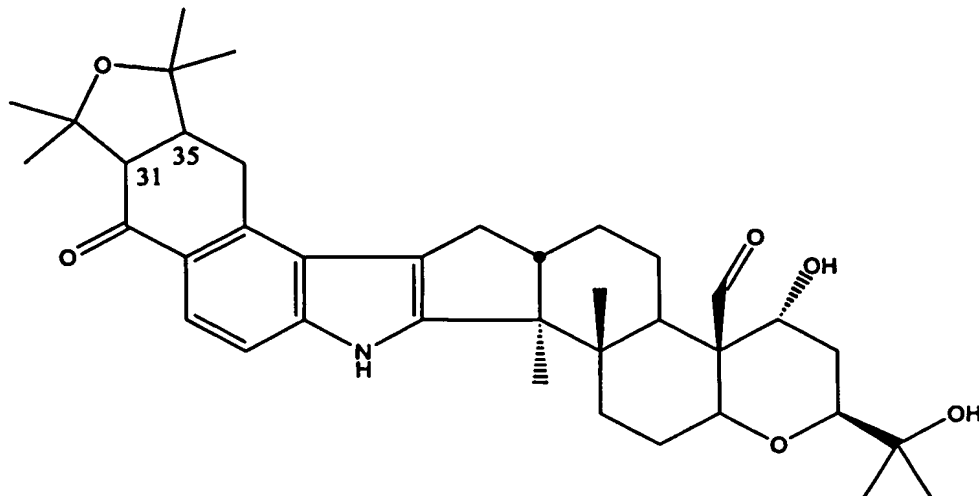
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STRUCTURE (X)

which includes lolicine A = 31 α , 35 β stereochemistry.

Preferably, the antagonist compound is structure (XI):



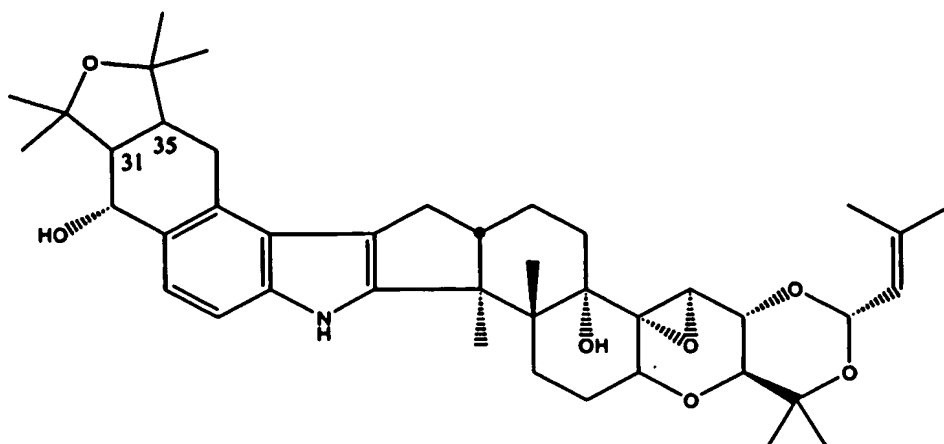
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STRUCTURE (XI)

which includes lolicine B = 31 α , 35 β stereochemistry.

Preferably, the antagonist compound is structure (XII):

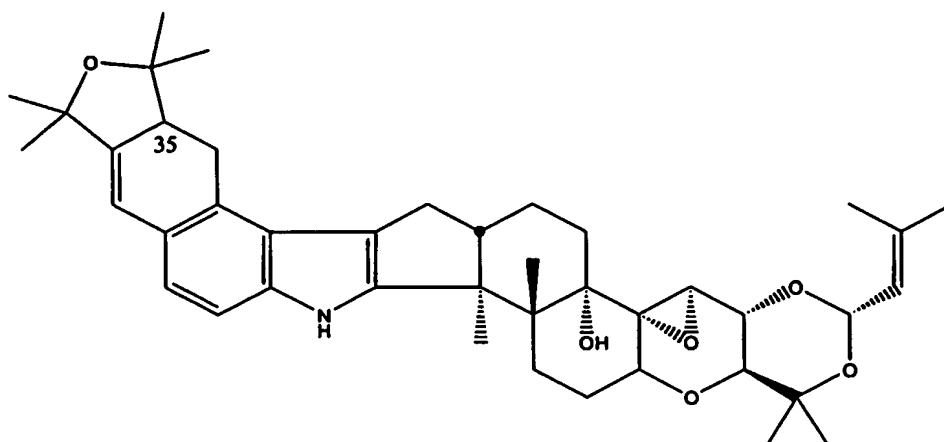
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STRUCTURE (XII)

which includes compounds selected from the group consisting of: 30-desoxylolitre B-30 α -ol = 31 α , 35 β stereochemistry; 30-desoxy-31-*epi*lolitre B-
 5 30 α -ol = 31 β , 35 β stereochemistry.

Preferably, the antagonist compound is structure (XIII):



STRUCTURE (XIII)

which includes 30-desoxylolitre B-30-ene = 35 β stereochemistry.

10 Preferably, the composition further includes pharmaceutically and physiologically acceptable carriers. Preferably, the pharmaceutically and physiologically

acceptable carriers include components selected from the group including; fillers; excipients; modifiers; humectants; stabilisers; emulsifiers; diluents; and other formulation components such as a use of a lipid vehicle.

Preferably, the composition, substantially as described above, is administered in a form selected from the group including: an injection; a tablet; a capsule; a suppository; an injection; a suspension; a drink or tonic; a syrup; a powder; an ingredient in solid or liquid foods; a nasal spray; a sublingual wafer; a transdermal patch; a transdermal injection; and combinations thereof. However, other methods of administration may also be employed without limiting the scope of the present invention.

Preferably, the BK channel antagonist compound or compounds are extracted from endophyte-infected plants and seeds; fungal cultures; chemical synthesis; heterologous expression systems including but not limited to bacteria, yeast, fungi, plants and animal cells; and combinations thereof.

In preferred embodiments, the source is perennial ryegrass seed from *Lolium perenne*. Further reference to lolitrem sources may be found in the applicants co-pending application NZ 530331.

Preferably, the ion channel is a potassium channel. More preferably, the potassium channel is a large conductance calcium activated potassium (BK) channel. Embodiments including intermediate conductance (IK) and small conductance (SK) may also be incorporated herein.

Preferably, the antagonist compound or compounds have activity against the alpha (α) subunit. More preferably, the antagonist compound or compounds have activity against both alpha (α) subunit and alpha plus beta (β) accessory subunit (β_1 to β_4) channels.

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Preferably, for lolitrem B, the degree of antagonist inhibition is approximately 97% for a composition containing approximately 20nM lolitrem B. The half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 3.7 ± 0.4 nM of lolitrem B.

- 5 Preferably, for lolitriol, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 1000 nM lolitriol. The half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 195 nM of lolitriol to inhibit α and β_1 activity and 536 ± 16 nM of lolitriol to inhibit α and β_4 activity.
- 10 Preferably, for 31-*epi*lolitrem B, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 200nM 31-*epi*lolitrem B. The half maximal degree of antagonist inhibition (IC_{50}) is found for a composition containing approximately 58 ± 6 nM of 31-*epi*lolitrem B to inhibit α and β_1 activity and 49 nM of 31-*epi*lolitrem B to inhibit α and β_4 activity.
- 15 Preferably, for lolitrem E, the degree of antagonist inhibition is approximately 100% for a composition containing approximately 100 nM lolitrem E.

It is the inventors understanding that the above levels of antagonist behaviour found in *in vitro* experiments indicates that lolitrem compounds have a high apparent affinity for at least *hSlo* channels. That is, the lolitrem compounds

20 reduce and inhibit potassium currents through *hSlo* channels as well as other *Slo* channels including but not limited to *mSlo* and *dSlo*.

The antagonist effect of the composition is not able to be reversed at larger concentrations by wash out for at least lolitrem B. It is inventors experience that this effect is for concentrations of 10 nM or greater of lolitrem B compound.

- 25 According to a further aspect of the present invention there is provided a method

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of preventing repolarisation or hyperpolarisation of a cell, wherein the cell contains a BK channel, including the administration to the cell of a pharmacologically effective amount of composition containing a BK channel antagonist substantially as described above.

- 5 According to a further aspect of the present invention there is provided the use of a composition substantially as described above for preventing repolarisation or hyperpolarisation of a cell, wherein the cell contains a BK channel.

It should be appreciated from the above description that there are provided compositions, methods and uses incorporating lolitrem compounds to antagonise
10 or block ion channel activity, particularly the BK channel.

It should further be appreciated that the blocking effect found for lolitrem compounds may be used in a variety of pharmaceutical applications such as for regulation of physiological functions including blood pressure, hypertension, muscle tone, brain functions such as neurotransmitter release and hearing
15 application. In addition, lolitrem compounds, and particularly lolitrem B, have been found to have a blocking effect that is strong and long acting. Further applications envisaged for this blocking effect includes use in drug development and diagnostics i.e. a known blocking effect is generated from lolitrem compounds which may be used to find new drugs or to test if various physiological effects are
20 altered from blocked channels. The above applications should not be seen as limiting as it should be appreciated by those skilled in the art that other applications may also be possible.

BRIEF DESCRIPTION OF THE DRAWINGS

Further aspects of the present invention will become apparent from the following description which is given by way of example only and with reference to the

5 accompanying drawings in which:

- Figure 1** shows two current recordings (A and B) for different concentrations of lolitrem B compared to control recordings of (A) ramp potential and (B) depolarising voltage pulses to +150 mV to determine the degree of antagonist inhibition;
- 10 **Figure 2** shows a graph of inhibition produced by different concentrations of lolitrem B. I/I_{max} is the current response as a fraction of the control response;
- Figure 3** shows two graphs (A and B) illustrating the effect of lolitrems on BK channel potassium currents for 31-*epi*lolitrem B;
- 15 **Figure 4** shows two graphs (A and B) illustrating the effect of lolitrems on BK channel potassium currents for lolitriol and channels containing different beta subunits; and
- Figure 5** shows a graph illustrating dose-response curves for macroscopic BK channel potassium currents inhibited by lolitrems. Current amplitude (I') 5 minutes after addition of increasing lolitriol concentrations, shown as a fraction of the control response. Lolitriol was applied to $\alpha+\beta_1$ channels (solid circles) and to $\alpha+\beta_4$ channels (open circles). 31-*epi*lolitrem B was applied to $\alpha+\beta_1$ channels (solid triangles) and to $\alpha+\beta_4$ channels (open triangles). Shaded triangles are data points from a single cell to which the curve was fitted.
- 20
- 25

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Lollitrem B was applied to hSlo channels (solid squares) a previous study and is shown for comparison. The current response was averaged over the last half of a voltage pulse to +150 mV for 50 ms, with 10 μ M internal free calcium. The vertical bars show \pm 1 S.E.M. in 3 or more cells. The curve is a fit of a Hill-type equation to the data.

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BEST MODES FOR CARRYING OUT THE INVENTION

The results found from experiments carried out by the inventors are now described.

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Experiment 1

In this experiment, hSlo α subunit large conductance calcium-activated potassium (BK) channels with an N-terminal c-myc tag in the mammalian vector pcDNA (Meera *et al*, 1997) were transiently expressed in human embryonic kidney cells (cell type HEK293).

15

Cell Culture Preparation

Human embryonic kidney cells were grown in a mix of DMEM (Dulbecco's Modified Eagle Medium, GibcoBRL Cat#12100-038) and 2.5 mM HEPES (N-[2-Hydroxyethyl]piperazine-N'-[2-ethanesulfonic acid]), supplemented with minimal essential amino acids and 10 % fetal bovine serum.

20

Cells were subsequently plated into 24-well plates, grown to 95% confluency and transfected 24 hours later with 10 µg of *hS/o* and 2 µg CD4 (pcDNA) and 2 µl Lipofectamine 2000™.

- Cells were plated onto cover slips 24 hours later. CD4 antibody-labelled beads
5 were used to identify transfected cells. Macroscopic currents were recorded from excised inside-out patches at 3 days post-transfection.

Lolitrems B preparation

Lolitrems B was extracted from perennial ryegrass seed infected with *Neotyphodium lolii*.

- 10 A stock of 100 µM Lolitrems B was made up in dimethyl sulfoxide (DMSO). This was diluted to the appropriate concentration in electrophysiological solutions. The final DMSO concentration was 0.1 % for 100 nM lolitrems B and did not exceed 0.02 % for lower concentrations.

Electrophysiology

- 15 *Solutions*

- The bath solution that was applied to the internal side of the cell membrane of the inside-out membrane patch was (mM): 140 KMeSO₃, 2 KCl, 20 HEPES (N-[2-Hydroxyethyl]piperazine-N'-[2-ethanesulfonic acid]), 5 HEDTA (N-[2-hydroxyethyl]ethylene-diaminetriacetic acid) and 3.65 CaCl₂ to give 10 µM free
20 calcium with a pH of 7.2.

The pipette solution applied to the external side of the cell membrane of the inside-out membrane patch was (mM): 140 KMeSO₃, 2 KCl, 20 HEPES, and 2 MgCl₂, with a pH of 7.2.

Macroscopic currents were recorded in an inside-out patch-clamp configuration using an amplifier, interface and data collection software. Data were filtered at 5 kHz and sampled at 20 μ s intervals. Fast capacitance compensation was used to cancel the fast transient. Leak subtraction was used although the background
5 potassium current was small.

Results

Inhibition of BK channels by lolitrem B

The effect of lolitrem B on potassium currents in excised inside-out patches from cells expressing *hSlo* BK channels is shown in Figure 1. The Figure shows two
10 current recordings (A and B) for different concentrations of lolitrem B compared to control recordings of (A) ramp potential and (B) depolarising voltage pulses to +150 mV to determine the degree of antagonist inhibition.

In Figure 2, solid circles show the mean normalised current (fraction blocked) in three or more cells and the vertical bars are mean ± 1 S.E.M. The curve is a fit of
15 the Hill equation to the data.

Figure 2 shows that the application of 20 nM lolitrem B to the perfusion bath, resulted in complete inhibition of the BK channel current. The level of antagonist inhibition was less at lower lolitrem B concentrations.

Channels were activated by voltage pulses to +150 mV every minute in the
20 presence of 10 μ M free calcium. Control current responses were recorded over 5 minutes. Only patches that remained stable over this time were used in experiments.

The current block produced by 10 nM or greater lolitrem B could not be reversed, even after wash-out with control solution for 30 minutes at a flow rate of 4ml/min in
25 three experiments.

At 2 nM lolitrem B partial inhibition was observed.

Dose-response experiments were carried out to determine the concentration range over which lolitrem B was effective.

Recovery from inhibition could not be used to validate reductions in current at

5 different lolitrem B concentrations as being due to the presence of drug, nor could different concentrations be applied in random order. However, by applying cumulative doses of lolitrem B to the same membrane patch, it was found that increases in the degree of current block with increased lolitrem B concentration were consistent between cells.

10 Each concentration of lolitrem B was applied for 5 minutes or until the current response had stabilised and fractional block calculated as the decrease in current as a fraction of the control. The current amplitude was the mean current over the last half of the voltage pulse to +150 mV. The data was analysed and fitted using the Hill equation which gave an estimate of the concentration of half maximal
15 inhibition (IC_{50}) of 3.7 ± 0.4 nM and a Hill coefficient of 1.7, from experiments using 11 cells.

The concentration range of inhibition observed for lolitrem B is similar to that reported for other indole diterpenes including: paxilline, aflatrem, penitrem A, paspalinine, paspalitrems A and C, verruculogen and paspalicine applied to BK
20 channels (Knaus et al., 1994, Sanchez and McManus, 1996, Gribkoff et al., 1996).

Difficulty in reversing channel block is also noted for these compounds, although paxilline block by low concentrations could be partially reversed by washout (Knaus et al., 1994).

Thus it can be seen from the above experiment that at least lolitrem B has a
25 blocking effect on at least BK channels.

Experiment 2

In this experiment, it is shown that whilst some lolitrem compounds are known to cause tremorgenicity to one extent or another, it is not certain that there is a direct
5 link to BK channel blocking and vice versa.

The experiment uses 31-*epi*lolitrem B, a known non-tremorgenic lolitrem compound as described in Munday-Finch et al 1996.

The same methods were used for testing the antagonist effect of lolitrem B as described in Experiment 1 above.

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Preparation of 31-*epi*lolitrem B

The lolitrem derivative 31-*epi*lolitrem B (Fig.1) was prepared by base-catalysed epimerization of lolitrem B according to S. Munday-Finch, 1997. The lolitrem B was extracted from ryegrass seed infected with *Neotyphodium lolii* (Gallagher et
15 al., 1981, Miles et al., 1994). A stock of 100 μ M 31-*epi*lolitrem B was made up in DMSO. This was diluted to the appropriate concentration in internal solution. The final DMSO concentration was 0.1 % for 100 nM compound and did not exceed 0.02 % for lower concentrations.

The results showed that 31-*epi*lolitrem B at a concentration of 100 nM inhibited BK
20 channel currents (α subunit) to 0.06 ± 0.02 (n=7) of the control response. The current response increased to 0.59 of the control response after wash-out with control solution for 26 minutes in one experiment,

Experiment 3

The experiment uses lolitriol, a known non-tremorgenic lolitrem compound as described in Munday-Finch, 1997.

The same methods were used for testing the antagonist effect of lolitrem B as
5 described in Experiment 1 above.

Preparation of lolitriol

The lolitrem derivative lolitriol was prepared by acid hydrolysis of lolitrem B as reported by Miles et al., 1992. The lolitrem B was extracted from ryegrass seed infected with *Neotyphodium lolii* (Gallagher et al., 1981, Miles et al., 1994). A
10 stock of 500 μ M lolitriol was made up in DMSO. This was diluted to the appropriate concentration in internal solution. The final DMSO concentration was 0.1 % for 1 μ M lolitriol and did not exceed 0.05 % for lower concentrations.

The results showed that lolitriol at a concentration of 100 nM inhibited BK channel currents (α subunit) to 0.29 (n=2) of the control response. 200 nM lolitriol inhibited
15 BK channel currents (α subunit) to 0.25 (n=1) of the control response. The current response increased to 0.86 of the control (n=2) response after wash-out with control solution for 30 minutes at a flow rate of 4ml/min in three experiments,

The results from Experiments 2 and 3 show that non-tremorgenic lolitrem compounds can inhibit BK channels. Tremorgenicity is thus unlikely to be directly
20 linked to BK channel blocking which is a similar result to that found in general indole diterpene studies (Knaus et al 1994).

Experiment 4

The experiment uses lolitrem E, a known partial-tremorgenic lolitrem compound as described in Munday-Finch, 1997. Lolitrem E is intermediate in structure between lolitrem B and lolitriol.

- 5 The same methods were used for testing the antagonist effect of lolitrem E as described in Experiment 1 above.

The results showed that lolitrem E at a concentration of 100 nM inhibited BK channel currents (α subunit) to 0.01 of the control response in one experiment.

10 **Experiment 5**

The aim of this experiment was to determine whether the non-tremorgenic lolitrems, 31-*epi*lolitrem B and lolitriol, inhibit function of BK channels that contain an accessory beta subunit. Assuming inhibition was found, it was also an aim to determine if inhibition is effected by the type of beta subunit present.

- 15 We used BK channels containing beta subunits that are expressed in smooth muscle (β_1) and brain (β_4). BK channels with subunit combinations $\alpha+\beta_1$ or $\alpha+\beta_4$ were expressed in human embryonic kidney cells and their function assayed by patch clamping.

Methods

- 20 *hSlo* α subunit large conductance calcium-activated potassium channels in pcDNA3 (Meera et al., 1997) together with the human β_4 subunit in pEGFP-N1 were transiently expressed in human embryonic kidney cells (HEK293). HEK cells were grown in a mix of DMEM (Dulbecco's Modified Eagle Medium) and 2.5 mM HEPES (N-[2-Hydroxyethyl]-piperazine-N'-[2-ethanesulfonic acid]), supplemented

with minimal essential amino acids and 10 % fetal bovine serum. Cells were subsequently plated onto 24-well plates, grown to 95% confluency and transfected 24 hours later with 10 µg of *hS/o*, 8 µg of β_4 and 2 µl Lipofectamine 2000™. Cells were plated onto cover slips 24 hours later. Transfected cells expressed green
5 fluorescent protein and were identified under UV light by their fluorescence.

Results

Both $\alpha+\beta_1$ and $\alpha+\beta_4$ BK channels produced potassium currents in response to depolarising voltages in the presence of 10 µM free calcium (Figure 4). The maximum current amplitude was typically 1-8 nA. The characteristics of the
10 current responses were similar to that previously reported for these channels (Brenner et al., 2000b, Ahring et al., 1997 Behrens et al., 2001 Lippiat et al., 2003)

31-*ep*lolitrem B and lolitriol inhibit BK channel function

Both 31-*ep*lolitrem B and lolitriol inhibited potassium currents through BK channels but at different concentrations (Figure 3 and Figure 4). The differences in the
15 concentration of the lolitrems required to inhibit BK channels is clearly shown in Figure 5. Dose response data for inhibition of BK channels by lolitrems are summarised below in Table 1. The IC₅₀ for inhibition by 31-*ep*lolitrem B was lower than that for lolitriol in both $\alpha+\beta_1$ (3 times lower) and $\alpha+\beta_4$ (10 times lower) BK channels. These results indicated that 31-*ep*lolitrem B has a higher apparent
20 affinity for BK channels than lolitriol in both $\alpha+\beta_1$ and $\alpha+\beta_4$ BK channels.

Table 1. Dose-response data for inhibition of BK channels by lolitrems

	<i>lolitrem B</i>		<i>31-epilolitrems B</i>		<i>lolitriol</i>	
	IC ₅₀	<i>h</i>	IC ₅₀	<i>h</i>	IC ₅₀	<i>h</i>
α alone	3.7 \pm 0.4 (n=11)	1.7	-	-	-	-
$\alpha+\beta_1$	-	-	58 \pm 6 (n=10)	2.7	195	2.3
$\alpha+\beta_4$	-	-	49 (n=5)	2.2	536 \pm 16 (n=7)	2.7

h = Hill coefficient

The above findings show that both tremorgenic and non-tremorgenic lolitrems can
 5 inhibit BK channel function. Therefore it is unlikely that BK channels mediate the
 tremorgenic actions of lolitrems, but rather that other molecular sites are involved.

A study of other indole diterpene compounds by McMillan et al., (2003) compared
 effects of tremorgenic versus non-tremorgenic compounds on BK channel function
 and also found differences between the two in their degree of block. The non-
 10 tremorgenic paxilline analogue, desoxypaxilline inhibits BK channels but requires
 24-times the concentration to produce the same degree of inhibition as for
 paxilline, which is tremorgenic.

Twice the concentration of lolitriol was required to inhibit $\alpha+\beta_4$ channels compared
 with $\alpha+\beta_1$ channels.

Experiment Summary

The above experiments determine the effect of four lolitrems (and by inference other related chemical structures): lolitrem B, lolitrem E, 31-*ep*lolitrem B and lolitriol on the function of BK channels. It is demonstrated that these compounds
5 inhibit potassium currents through BK channels.

The results also show that BK channels that contain accessory beta subunits are also inhibited by lolitrem B, lolitrem E, 31-*ep*lolitrem B and lolitriol.

It is envisaged by the inventors that, because 31-*ep*lolitrem B and lolitriol inhibit BK channel function and are non-tremorgenic in mice, they may have potential as
10 research tools in the study of BK channel pharmacology or as drugs.

The relatively low concentration of 31-*ep*lolitrem B, that is sufficient for BK inhibition, together with its non-tremorgenic properties, suggest uses for this lolitrem derivative in *in vivo* applications where a BK channel blocker is required.

Aspects of the present invention have been described by way of example only and
15 it should be appreciated that modifications and additions may be made thereto without departing from the scope thereof as defined in the appended claims.

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